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REQUEST **FOR** 

## **CONTINUED EXAMINATION (RCE)** TRANSMITTAL

Subsection (b) of 35 U.S.C. §132, effective on May 29, 2000, provides for continued examination of a utility or plant application filed on or after June 8,1995.

See the American Inventors Protection Act of 1999 (AIPA).

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Application Number	09/782,184		
Filing Date	February 12, 200		25
Tilling Date	1 0010017 12, 200	<del>~</del>	$\leq$
First Named Inventor	Sands	<u> </u>	-
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Group Art Unit	1616	<u>ன்</u>	407
		$\aleph^-$	$\sim$
Examiner Name	Gollamundi	<u> </u>	
		8	
Attorney Docket Number	12636-898		

This is a Request for Continued Examination (RCE) under 37 C.F.R. §1.114 of the above-identified application.

NOTE: 37 C.F.R. § 1.114 is effective on May 29, 2000. If the above-identified application was filed prior to May 29, 2000, applicant may wish to consider filing a continued prosecution application (CPA) under 37 C.F.R. s 1.53(d) (PTO/SB/29) instead of a RCE to be eligible for the patent term adjustment provisions of the AIPA. See Changes to Application Examination and Provisional Application Practice, Interim Rule, 65 Fed. Reg. 14865 (Mar. 20, 2000), 1233 Off. Gaz. Pat. Office 47 (Apr. 11, 20000), which established RCE practice.

1.	Sub	missio	n required under 37 C.F.R. §1.114.					
	a.	. Previously submitted						
		i.	Consider the amendment(s)/reply under 37 C.F.R. §1.116 previously filed on (Any unentered amendment(s) referred to above will be entered)					
	ii. Consider the arguments in the Appeal Brief or Reply Brief previously filed on							
		iii.	Other	ļ				
	b.	$\boxtimes$	Enclosed .					
		i.	□ Preliminary Amendment/Reply					
		ii.	Affidavit(s)/Declaration(s)					
	iii.   Information Disclosure Statement (IDS) (1 Reference Cited)							
		iv.	☐ Other	.				
2.	Mic	cellan	anus					
۷.			Suspension of action on the above-identified application is requested under 37 C.F.R. § 1.103(c) for a					
	a.	ш	period of months. (Period of suspension shall not exceed 3 months; Fee under 37 C.F. R. § 1.17(i) required)					
	b.		Other					
3.	Fees The RCE fee under 37 C.F.R. §1.17(e) is required by 37 C.F.R. § 1.114 when the RCE is filed.							
	a.   The Director is hereby authorized to charge the following fees, or credit any overpayments to Deposit							
			Account No. 23-2415 (Docket No. 12636-898)					
		i.	RCE fee required under 37 C.F.R. § 1.17(e)					
		ii.	Extension of time fee (37 C.F.R. §§ 1.136 and 1.17) (2 Month Extension)  11/18/2002 ANDNDAF1 00000173 232415 09782184					
		iii.	Other state of the					
	b.		Check in the amount of \$ enclosed 01 FC:1801 740.00 CH					
	C.		Payment by credit card (Form PTO-2038 enclosed) 02 FC:1252 400.00 CH					
				$\neg$				
SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT REQUIRED								
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Sigi	nature		Mate   November 13, 2002					
	-		CERTIFICATE OF MAILING OR TRANSMISSION	$\neg$				
I he	eby ce	ertify tha	Abic surrections is being deposited with the United States Postal Service as first class mail in an envelope addressed to:					
Con 200	nmissio	oner for	t this correspondence is being deposited with the Office of States Foods of States and Trademark Office on: November 13 Patents, Box RCE, Washington, DC 20231, or facsimile transmitted to the U.S. Patent and Trademark Office on: November 13					
Nar		Print/Typ	Annette Parladino					
Signature Lunttalladus Date November 13, 2002								
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11/19/05 11/19/05

Practitioner's Docket No. 12636-898

IN T	HE UNITED STATES PATEN	T AND TRADEMARK OFFIC	
In re application	of: Howard Sands, et al.	)	CEIVED 1
Serial No.: 09/	782,184	) Group No.: 1616	
Filed: February	12, 2001	) Examiner: Gollamudi, Sh	armila S.
Compos	ole Pharmaceutical sition Comprising roplets of A Camptothecin	) ) )	

Commissioner for Patents Washington, D.C. 20231

## PRELIMINARY AMENDMENT

Sir:

Prior to an examination on the merits of the above-identified application please amend the application as follows:

## In the Claims:

Please amend independent claims 1, 18-19, and dependent claims 22-23 as follows:

1. (Twice Amended) An injectable pharmaceutical composition comprising: an aqueous suspension of microdroplets suitable for intravenous delivery, the microdroplets having a mean diameter between 200 Angstroms and one micron, the microdroplets comprising a substantially water-insoluble, pharmacologically acceptable liquid that does not tend to form micelle structures, a camptothecin dissolved in water-insoluble, and pharmacologically acceptable liquid, and an outer layer comprising a

phospholipid.

